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VAR G1=8/12
REP G2=(0-10) C
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CONNECT IS E1 RC AT 13
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

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NUMBER OF NODES IS 13

STEREO ATTRIBUTES: NONE

DIENEO	WIINTDOIL	DD. MOME			
L3	98	SEA FILE=REGISTE	RY SSS FU	L L1	·
L4	21	SEA FILE=CAPLUS	ABB=ON	PLU=ON	L3
L5	242	SEA FILE=CAPLUS	ABB=ON	PLU=ON	("IHARA H"/AU OR "IHARA
		HIDEAKI"/AU OR '	'IHARA HI	DEKI"/AU	J)
L6	51	SEA FILE=CAPLUS	ABB=ON	PLU=ON	("TAKAOKA D"/AU OR "TAKAOKA
		DAISUKE"/AU)			
L7	186	SEA FILE=CAPLUS	ABB=ON	PLU=ON	("MIZUNO H"/AU OR "MIZUNO
		HAJIME"/AU)			
L8	474	SEA FILE=CAPLUS	ABB=ON	PLU=ON	(L5 OR L6 OR L7)
L9	6	SEA FILE=CAPLUS	ABB=ON	PLU=ON	L8 AND THIA? AND ?AZOL?
L10	1	SEA FILE=CAPLUS	ABB=ON	PLU=ON	L9 AND L4

=> d 112 ibib abs hitstr tot

L12 ANSWER 1 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN ACCESSION NUMBER: 2006:680917 CAPLUS Full-text

DOCUMENT NUMBER: 145:145750

TITLE: Preparation of pyrrolidine derivatives as

dipeptidylpeptidase IV inhibitors

INVENTOR(S): Nakai, Hisao; Kondo, Takashi; Ota, Motohiro

PATENT ASSIGNEE(S): Ono Pharmaceutical Co., Ltd., Japan

SOURCE: PCT Int. Appl., 163 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

P.	ATENT 1	NO.			KINI	D 1	DATE		1	APPL.	ICAT.	LONI	NO. 		D <i>I</i>	ATE 	
WC	2006	0731	 67		A1		2006	0713	. 7	WO 20	006-	JP30	0061		20	00601	L06
	W:	ΑE,	AG,	AL,	AM,	AT,	AU,	AZ,	BA,	BB,	BG,	BR,	BW,	BY,	ΒZ,	CA,	CH,
		CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GD,
		GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KM,	KN,	ΚP,	KR,
																MW,	
																SD,	
		SG,	SK,	SL,	SM,	SY,	ТJ,	TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VC,
					ZM,												
	RW:															HU,	
		IS,	IT,	LT,	LU,	LV,	MC,	NL,	PL,	PT,	RO,	SE,	SI,	SK,	TR,	BF,	ВJ,
																BW,	
		GM,	KE,	LS,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	ΑZ,	BY,
		KG,	ΚZ,	MD,	RU,	ТJ,	$\mathbf{M}\mathbf{T}$										
PRIORI	TY APP	LN.	INFO	.:						JP 2	005-	3063			A 2	0050	107
OTHER :	SOURCE	(S):			MAR	PAT	145:	1457	50								

$$Z=Y \xrightarrow{\left(\begin{array}{c} B \\ \end{array}\right)} W \xrightarrow{\left(\begin{array}{c} \left(R11\right) p \\ \end{array}} W \xrightarrow{\left(\begin{array}{c} \left(R11\right) p \\ \end{array}\right)} W \xrightarrow{\left(\begin{array}{c} \left(R11\right) p \\ \end{array}} W \xrightarrow{\left(\begin{array}{c}$$

The title compds. I [V, W and Y represent each a bond or a spacer having from 1 to 8 atoms in the main chain; the rings A and B are each a cyclic group optionally further having substituent(s); Z represents H or a substituent; X represents carbon or sulfur; R11 and R12 represent each a substituent; p and q are each 0 or an integer of 1 to 4; and x and m are each 0 or 1; the dotted line indicates a single bond or a double bond; α and β or β and γ do not

represent double bonds at the same time; when X is S, both α and β indicate single bonds] are prepared Thus, 1-(3-methyl-1,2,4-thiadiazol-5-yl)-4-([(3S,5S)-5- (pyrrolidin-1-ylcarbonyl)pyrrolidin-3-yl]carbonyl)piperazine hydrochloride was prepared in a multistep process from 2-benzyl 1-tert-Bu (2S,4S)-4-cyano-1,2-pyrrolidinedicarboxylate. Compds. of this invention showed IC50 values of 18 nM to 52 nM against dipeptidylpeptidase IV. Formulations are given.

IT 898274-42-5P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of pyrrolidine derivs. as dipeptidylpeptidase IV inhibitors)

RN 898274-42-5 CAPLUS

CN Piperazine, 1-[3-(methylthio)-1,2,4-thiadiazol-5-yl]-4-[[(3S,5S)-5-(1-pyrrolidinylcarbonyl)-3-pyrrolidinyl]carbonyl]-, monohydrochloride (9CI) (CA INDEX NAME)

Absolute stereochemistry.

$$\begin{array}{c|c} & & & & \\ & &$$

● HCl

REFERENCE COUNT:

THERE ARE 58 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L12 ANSWER 2 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN

58

ACCESSION NUMBER:

2006:489438 CAPLUS Full-text

DOCUMENT NUMBER:

144:462620

TITLE:

SOURCE:

Pest control compositions containing imidacloprid and

pyrimidines or thiadiazoles

INVENTOR(S):

Shimokawadoko, Yasutaka; Yamada, Koji Sumitomo Chemical Co., Ltd., Japan

PATENT ASSIGNEE(S):

Jpn. Kokai Tokkyo Koho, 46 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent Japanese

LANGUAGE:

Japan

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2006131536	Α	20060525	JP 2004-321494	20041105
PRIORITY APPLN. INFO.:			JP 2004-321494	20041105
OTHER SOURCE(S):	MARPAT	144:462620		
GI				

$$X_1$$
 X_2 X_3 X_4 X_4 X_5 X_5 X_6 X_7 X_8 X_8 X_8 X_8 X_8 X_8 X_8 X_8 X_8 X_8

AB Pest control compns. with excellent efficacy contain (1) a pyrimidine compound (I, wherein Rl = H, alkyl; R2 = alkynyloxy; R3 = H, halo, alkyl; X1 = (un)substituted C4-7 polymethylene) or a thiadiazole and (2) imidacloprid as active components. Thus, 4-(2-butynyloxy)-5-fluoro-6-(3,3-dimethylpyrrolidin-1-yl)pyrimidine + Admire wettable powder at 12.5 + 0.025 ppm was more effective than the individual components at the same rates against Aphis gossypii on cucumber.

IT 886841-46-9 886841-48-1 886841-50-5
RL: AGR (Agricultural use); BSU (Biological study, unclassified); BIOL (Biological study); USES (Uses)
(as pesticide)

RN 886841-46-9 CAPLUS

CN 2-Imidazolidinimine, 1-[(6-chloro-3-pyridinyl)methyl]-N-nitro-, (2E)-, mixt. with 3-methyl-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]piperidine (9CI) (CA INDEX NAME)

CM 1

CRN 850748-38-8 CMF C13 H19 N3 O S

$$Et-C = C-CH_2-O$$
 $N-S$
 $N-S$

CM 2

CRN 138261-41-3 CMF C9 H10 C1 N5 O2

RN 886841-48-1 CAPLUS

CN 2-Imidazolidinimine, 1-[(6-chloro-3-pyridinyl)methyl]-N-nitro-, (2E)-, mixt. with 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-dimethylpiperidine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-59-1 CMF C13 H19 N3 O S

$$Me-C = C-CH_2-O$$
 N
 N
 Me
 Me
 Me

CM 2

CRN 138261-41-3 CMF C9 H10 C1 N5 O2

RN 886841-50-5 CAPLUS

CN 2-Imidazolidinimine, 1-[(6-chloro-3-pyridinyl)methyl]-N-nitro-, (2E)-, mixt. with 2,6-dimethyl-4-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]morpholine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-61-5 CMF C13 H19 N3 O2 S

$$\mathsf{Et-C} = \mathsf{C-CH_2-O} \overset{\mathsf{Me}}{\underset{\mathsf{N-S}}{\overset{\mathsf{Me}}{\longrightarrow}}} \overset{\mathsf{Me}}{\underset{\mathsf{Me}}{\overset{\mathsf{O}}{\longrightarrow}}}$$

CRN 138261-41-3 CMF C9 H10 C1 N5 O2

L12 ANSWER 3 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2006:489434 CAPLUS Full-text

DOCUMENT NUMBER:

144:462619

TITLE:

Pest control compositions containing acetamiprid and

pyrimidines or thiadiazoles

INVENTOR(S):

Shimokawadoko, Yasutaka; Yamada, Koji Sumitomo Chemical Co., Ltd., Japan

PATENT ASSIGNEE(S): SOURCE:

Jpn. Kokai Tokkyo Koho, 47 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

GI

Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2006131537	Α	20060525	JP 2004-321495	20041105
PRIORITY APPLN. INFO.:			JP 2004-321495	20041105
OTHER SOURCE(S):	MARPAT	144:462619		

$$X_1$$
 X_2 X_3 X_4 X_4 X_5 X_5

AB Highly effective pest control compns. contain (1) pyrimidines (I, wherein R1 = H, alkyl; R2 = alkynyloxy; R3 = H, halo, alkyl; X1 = (un)substituted C4-7 polymethylene) or 1,2,4-thiadiazole derivs. such as 3-(2-pentynyloxy)-5-(3-methylpiperidino)-1,2,4-thiadiazole and (2) (E)-N1-[(6-chloro-3-pyridyl)methyl]-N2-cyano-N1-methylacetamidine (acetamiprid). Thus, 4-(2-

butynyloxy)-5-fluoro-6-(3,3-dimethylpyrrolidin-1-yl)pyrimidine + acetamiprid at 12.5 + 0.025 ppm was more effective than the individual components at the same rates against Aphis gossypii on cucumber.

IT 886840-72-8 886840-73-9 886840-74-0

RL: AGR (Agricultural use); BSU (Biological study, unclassified); BIOL (Biological study); USES (Uses)

(as pesticide) 886840-72-8 CAPLUS

CN Ethanimidamide, N-[(6-chloro-3-pyridinyl)methyl]-N'-cyano-N-methyl-, (1E)-, mixt. with 3-methyl-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]piperidine (9CI) (CA INDEX NAME)

CM 1

RN

CRN 850748-38-8 CMF C13 H19 N3 O S

$$Et-C = C-CH_2-O$$
 $N-S$
 $M \in S$

CM 2

CRN 135410-20-7 CMF C10 H11 C1 N4

Double bond geometry as shown.

RN 886840-73-9 CAPLUS

CN Ethanimidamide, N-[(6-chloro-3-pyridinyl)methyl]-N'-cyano-N-methyl-, (1E)-, mixt. with 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-dimethylpiperidine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-59-1 CMF C13 H19 N3 O S

$$Me-C = C-CH_2-O$$
 $N-S$
 Me
 Me
 Me

CRN 135410-20-7 CMF C10 H11 C1 N4

Double bond geometry as shown.

RN 886840-74-0 CAPLUS

CN Ethanimidamide, N-[(6-chloro-3-pyridinyl)methyl]-N'-cyano-N-methyl-, (1E)-, mixt. with 2,6-dimethyl-4-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]morpholine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-61-5 CMF C13 H19 N3 O2 S

CM 2

CRN 135410-20-7 CMF C10 H11 C1 N4

Double bond geometry as shown.

ACCESSION NUMBER:

2006:489433 CAPLUS Full-text

DOCUMENT NUMBER:

144:462618

TITLE:

Pest control compositions containing pyriproxyfen and

pyrimidines or thiadiazoles

INVENTOR(S):
PATENT ASSIGNEE(S):

Shimokawadoko, Yasutaka; Yamada, Koji Sumitomo Chemical Co., Ltd., Japan

SOURCE:

LANGUAGE:

Jpn. Kokai Tokkyo Koho, 46 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2006131528	Α	20060525	JP 2004-321486	20041105
PRIORITY APPLN. INFO.:		·	JP 2004-321486	20041105
OTHER SOURCE(S):	MARPAT	144:462618		
GI				

AB Highly effective pest control compns. contain (1) pyrimidines (I, wherein R1 = H, alkyl; R2 = alkynyloxy; R3 = H, halo, alkyl; X1 = (un)substituted C4-7 polymethylene) or 1,2,4-thiadiazole derivs. such as 3-(2-pentynyloxy)-5-(3-methylpiperidino)-1,2,4-thiadiazole and (2) 4-phenoxyphenyl (RS)-2-(2-pyridyloxy) Pr ether (pyriproxyfen) as active components. Thus, 4-(2-butynyloxy)-5-fluoro-6-(3,3-dimethylpyrrolidin-1-yl)pyrimidine + pyriproxyfen at 0.15 + 50 ppm showed greater control of Bemisia argentifolii on cabbage than the individual components at the same resp. rates.

IT 886840-29-5 886840-30-8 886840-31-9

RL: AGR (Agricultural use); BSU (Biological study, unclassified); BIOL (Biological study); USES (Uses)

(as pesticide)

RN 886840-29-5 CAPLUS

CN Pyridine, 2-[1-methyl-2-(4-phenoxyphenoxy)ethoxy]-, mixt. with 3-methyl-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]piperidine (9CI) (CA INDEX NAME)

CM 1

CRN 850748-38-8 CMF C13 H19 N3 O S

CRN 95737-68-1 CMF C20 H19 N O3

RN 886840-30-8 CAPLUS

CN Pyridine, 2-[1-methyl-2-(4-phenoxyphenoxy)ethoxy]-, mixt. with 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-dimethylpiperidine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-59-1 CMF C13 H19 N3 O S

$$Me = C = C + 2 = 0$$
 $N = S$
 $N = S$
 $N = S$
 $N = S$

CM 2

CRN 95737-68-1 CMF C20 H19 N O3

RN 886840-31-9 CAPLUS

CN Morpholine, 2,6-dimethyl-4-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]-, mixt. with 2-[1-methyl-2-(4-phenoxyphenoxy)ethoxy]pyridine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-61-5 CMF C13 H19 N3 O2 S

CM 2

CRN 95737-68-1 CMF C20 H19 N O3

L12 ANSWER 5 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2006:489432 CAPLUS Full-text

DOCUMENT NUMBER:

144:462617

TITLE:

Pest control compositions containing thiamethoxam and

pyrimidines or thiadiazoles

INVENTOR(S):

Shimokawadoko, Yasutaka; Yamada, Koji Sumitomo Chemical Co., Ltd., Japan

PATENT ASSIGNEE(S):

SOURCE:

Jpn. Kokai Tokkyo Koho, 47 pp.

DOCUMENT TYPE:

CODEN: JKXXAF Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2006131530 PRIORITY APPLN. INFO.:	Α	20060525	JP 2004-321488 JP 2004-321488	20041105 20041105
OTHER SOURCE(S):	MARPAT	144:462617		

$$Et-C = C-CH_2-O \xrightarrow{N-S} N$$
II

- AB Compns. with excellent pest control effect contain (1) a pyrimidine compound (e.g., I) or a 1,2,4-thiadiazole derivative (e.g., II) and (2) 3-[(2-chloro-1,3-thiazol-5-yl)methyl]-5-methyl-1,3,5-oxadiazinan-4- ylidene(nitro)amine (thiamethoxam). Thus, I + thiamethoxam at 12.5 + 0.025 ppm was more effective than the individual components at the same rates against Aphis gossypii on cucumber.
- RN 886843-19-2 CAPLUS
- CN 4H-1,3,5-Oxadiazin-4-imine, 3-[(2-chloro-5-thiazolyl)methyl]tetrahydro-5-methyl-N-nitro-, mixt. with 3-methyl-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]piperidine (9CI) (CA INDEX NAME)

CRN 850748-38-8 CMF C13 H19 N3 O S

CM 2

CRN 153719-23-4 CMF C8 H10 C1 N5 O3 S

$$\begin{array}{c|c}
 & \text{Me} \\
 & \text{N} & \text{NO2} \\
 & \text{N} & \text{CH}_2 & \text{S} \\
 & \text{N} & \text{C1}
\end{array}$$

RN 886843-21-6 CAPLUS

CN 4H-1,3,5-Oxadiazin-4-imine, 3-[(2-chloro-5-thiazolyl)methyl]tetrahydro-5-methyl-N-nitro-, mixt. with 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-dimethylpiperidine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-59-1 CMF C13 H19 N3 O S

$$Me-C = C-CH_2-O$$
 $N-S$
 Me
 Me
 Me

CM 2

CRN 153719-23-4 CMF C8 H10 C1 N5 O3 S

RN 886843-22-7 CAPLUS

CN 4H-1,3,5-Oxadiazin-4-imine, 3-[(2-chloro-5-thiazolyl)methyl]tetrahydro-5-methyl-N-nitro-, mixt. with 2,6-dimethyl-4-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]morpholine (9CI) (CA INDEX NAME)

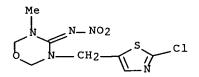
CM 1

CRN 886757-61-5 CMF C13 H19 N3 O2 S

$$Et-C = C-CH_2-O N O N$$

CM 2

CRN 153719-23-4 CMF C8 H10 C1 N5 O3 S



L12 ANSWER 6 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2006:489274 CAPLUS Full-text

DOCUMENT NUMBER: 144:462616

TITLE: Pest control compositions containing fenpropathrin and

pyrimidines or thiadiazoles

INVENTOR(S): Shimokawatoko, Yasutaka; Yamada, Koji

PATENT ASSIGNEE(S): Sumitomo Chemical Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 44 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

IT

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2006131541 PRIORITY APPLN. INFO.:	Α	20060525	JP 2004-321499 JP 2004-321499	20041105 20041105
OTHER SOURCE(S):	MARPAT	144:462616		

AB Compns. with excellent pest control effect contain (1) a pyrimidine derivative (e.g., I) or a 1,2,4-thiadiazole (e.g., II) and (2) fenpropathrin. Thus, 4-(2-butynyloxy)-5-fluoro-6-(3,3-dimethylpyrrolidin-1-yl)pyrimidine + Rody (500 + 100 ppm) gave almost 100% control of Aphis gossypii on cucumber.

886974-51-2 886974-52-3 886974-53-4
RL: AGR (Agricultural use); BSU (Biological study, unclassified); BIOL (Biological study); USES (Uses)
(as pesticide)

RN 886974-51-2 CAPLUS

CN Cyclopropanecarboxylic acid, 2,2,3,3-tetramethyl-, cyano(3-phenoxyphenyl)methyl ester, mixt. with 3-methyl-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]piperidine (9CI) (CA INDEX NAME)

CM 1

CRN 850748-38-8 CMF C13 H19 N3 O S

$$Et-C = C-CH_2-O N N-S N$$

CM 2

CRN 39515-41-8 CMF C22 H23 N O3

RN 886974-52-3 CAPLUS

CN Cyclopropanecarboxylic acid, 2,2,3,3-tetramethyl-, cyano(3-phenoxyphenyl)methyl ester, mixt. with 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-dimethylpiperidine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-59-1 CMF C13 H19 N3 O S

$$Me-C = C-CH_2-O$$
 $N-S$
 Me
 Me
 Me

CM 2

CRN 39515-41-8 CMF C22 H23 N O3

$$\begin{array}{c} \text{Me} & \text{Me} \\ \text{Me} & \text{C-O-CH} \\ \text{Ne} & \text{CN} \end{array}$$

RN886974-53-4 CAPLUS

CN Cyclopropanecarboxylic acid, 2,2,3,3-tetramethyl-, cyano(3phenoxyphenyl)methyl ester, mixt. with 2,6-dimethyl-4-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]morpholine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-61-5 CMF C13 H19 N3 O2 S

$$Et-C = C-CH_2-O \xrightarrow{N} \xrightarrow{Me} O$$

2 CM

39515-41-8 CRN CMF C22 H23 N O3

L12 ANSWER 7 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN ACCESSION NUMBER: 2006:489273 CAPLUS Full-text

DOCUMENT NUMBER: 144:462615

TITLE: Pest control compositions containing

 λ -cyhalothrin and pyrimidines or thiadiazoles

INVENTOR(S): Shimokawatoko, Yasutaka; Yamada, Koji

PATENT ASSIGNEE(S): Sumitomo Chemical Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 45 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
		-		-
JP 2006131539 PRIORITY APPLN. INFO.:	Α	20060525	JP 2004-321497 JP 2004-321497	20041105 20041105
OTHER SOURCE(S):	MARPAT	144:462615		

AB Compns. with excellent efficacy in pest control contain (1) a pyrimidine (I, wherein Rl = H, alkyl; R2 = alkynyloxy; R3 = H, halo, alkyl; X1 = (un)substituted C4-7 polymethylene) or a thiadiazole compound such as $3-(2-pentynyloxy)-5-(3-methylpiperidino)-1,2,4-thiadiazole and (2) <math>\lambda$ -cyhalothrin. Thus, 4-(2-butynyloxy)-5-fluoro-6-(3,3-dimethylpyrrolidin-1-yl)pyrimidine + Karate at 12.5 + 1.5 ppm was more effective than the individual components at the same rates against Aphis gossypii on cucumber.

IT 886844-06-0 886844-07-1 886844-08-2
RL: AGR (Agricultural use); BSU (Biological study, unclassified); BIOL (Biological study); USES (Uses)
(as pesticide)

RN 886844-06-0 CAPLUS

CN Cyclopropanecarboxylic acid, 3-[(1Z)-2-chloro-3,3,3-trifluoro-1-propenyl]-2,2-dimethyl-, (R)-cyano(3-phenoxyphenyl)methyl ester, (1S,3S)-rel-, mixt. with 3-methyl-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]piperidine (9CI) (CA INDEX NAME)

CM 1

CRN 850748-38-8 CMF C13 H19 N3 O S

$$Et-C = C-CH_2-O$$
 $N-S$
 Me

CM 2

CRN 91465-08-6 CMF C23 H19 C1 F3 N O3

Relative stereochemistry.

Double bond geometry as shown.

RN 886844-07-1 CAPLUS

CN Cyclopropanecarboxylic acid, 3-[(12)-2-chloro-3,3,3-trifluoro-1-propenyl]-2,2-dimethyl-, (R)-cyano(3-phenoxyphenyl)methyl ester, (1S,3S)-rel-, mixt. with 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-dimethylpiperidine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-59-1 CMF C13 H19 N3 O S

CM 2

CRN 91465-08-6 CMF C23 H19 Cl F3 N O3

Relative stereochemistry.

Double bond geometry as shown.

RN 886844-08-2 CAPLUS

CN Cyclopropanecarboxylic acid, 3-[(1Z)-2-chloro-3,3,3-trifluoro-1-propenyl]-2,2-dimethyl-, (R)-cyano(3-phenoxyphenyl)methyl ester, (1S,3S)-rel-, mixt. with 2,6-dimethyl-4-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]morpholine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-61-5 CMF C13 H19 N3 O2 S

CRN 91465-08-6 CMF C23 H19 C1 F3 N O3

Relative stereochemistry.

Double bond geometry as shown.

L12 ANSWER 8 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2006:489271 CAPLUS Full-text

DOCUMENT NUMBER:

144:462614

TITLE:

Pest control compositions containing deltamethrin and

pyrimidines or thiadiazoles

INVENTOR(S):

Shimokawatoko, Yasutaka; Yamada, Koji

PATENT ASSIGNEE(S):

Sumitomo Chemical Co., Ltd., Japan

SOURCE:

Jpn. Kokai Tokkyo Koho, 45 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2006131538	Α	20060525	JP 2004-321496	20041105
PRIORITY APPLN. INFO.:			JP 2004-321496	20041105
OTHER SOURCE(S):	MARPAT	144:462614		
GI				

Pest control compns. with superior efficacy contain (1) a pyrimidine (I, wherein R1 = H, alkyl; R2 = alkynyloxy; R3 = H, halo, alkyl; X1 = (un)substituted C4-7 polymethylene) or a 1,2,4-thiadiazole derivative such as 3-(2-pentynyloxy)-5-(3-methylpiperidino)-1,2,4-thiadiazole and (2) deltamethrin. Thus, 4-(2-butynyloxy)-5-fluoro-6-(3,3-dimethylpyrrolidin-1-yl)pyrimidine + deltamethrin emulsion (Decis) at 12.5+ 1.5 ppm was more effective than the individual components at the same rates against Aphis gossypii on cucumber.

IT 886841-88-9 886841-90-3 886841-92-5

RL: AGR (Agricultural use); BSU (Biological study, unclassified); BIOL (Biological study); USES (Uses)

(as pesticide)

RN 886841-88-9 CAPLUS

CN Cyclopropanecarboxylic acid, 3-(2,2-dibromoethenyl)-2,2-dimethyl-, (S)-cyano(3-phenoxyphenyl)methyl ester, (1R,3R)-, mixt. with 3-methyl-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]piperidine (9CI) (CA INDEX NAME)

CM 1

CRN 850748-38-8 CMF C13 H19 N3 O S

$$Et-C = C-CH_2-O$$
 $N-S$
 $N-S$
 $N-S$

CM 2

CRN 52918-63-5

CMF C22 H19 Br2 N O3

Absolute stereochemistry.

RN 886841-90-3 CAPLUS

CN Cyclopropanecarboxylic acid, 3-(2,2-dibromoethenyl)-2,2-dimethyl-, (S)-cyano(3-phenoxyphenyl)methyl ester, (1R,3R)-, mixt. with 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-dimethylpiperidine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-59-1

CMF C13 H19 N3 O S

$$Me-C = C-CH_2-O$$
 $N-S$
 Me
 Me
 Me

CM 2

CRN 52918-63-5

CMF C22 H19 Br2 N O3

Absolute stereochemistry.

RN 886841-92-5 CAPLUS

CN Cyclopropanecarboxylic acid, 3-(2,2-dibromoethenyl)-2,2-dimethyl-, (S)-cyano(3-phenoxyphenyl)methyl ester, (1R,3R)-, mixt. with 2,6-dimethyl-4-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]morpholine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-61-5 CMF C13 H19 N3 O2 S

CM 2

CRN 52918-63-5

CMF C22 H19 Br2 N O3

Absolute stereochemistry.

L12 ANSWER 9 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2006:489270 CAPLUS Full-text

DOCUMENT NUMBER: 144:462613

TITLE: Pest control compositions containing dinotefuran and

pyrimidines or thiadiazoles

INVENTOR(S): Shimokawatoko, Yasutaka; Yamada, Koji PATENT ASSIGNEE(S): Sumitomo Chemical Co., Ltd., Japan

PATENT ASSIGNEE(S): Sumitomo Chemical Co., Ltd., Japan SOURCE: Jpn. Kokai Tokkyo Koho, 46 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2006131533	Α	20060525	JP 2004-321491	20041105
PRIORITY APPLN. INFO.:			JP 2004-321491	20041105
OTHER SOURCE(S):	MARPAT	144:462613		
GI				

Me
$$\stackrel{\text{Me}}{\longrightarrow}$$
 N $\stackrel{\text{N}}{\longrightarrow}$ N $\stackrel{\text{O-CH}_2-C}{\equiv}$ C $\stackrel{\text{CH}_3}{\longrightarrow}$ I

AB Pest control compns. with superior efficacy contain (1) a pyrimidine compound (e.g., I) or a 1,2,4-thiadiazole derivative (e.g., II) and (2) dinotefuran. Thus, I + dinotefuran at 12.5 + 0.025 ppm was more effective than the individual components at the same rates against Aphis gossypii on cucumber.

RN 886757-58-0 CAPLUS

CN Guanidine, N-methyl-N'-nitro-N''-[(tetrahydro-3-furanyl)methyl]-, mixt. with 3-methyl-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]piperidine (9CI) (CA INDEX NAME)

CRN 850748-38-8 CMF C13 H19 N3 O S

$$Et-C = C-CH_2-O N N-S N-S N$$

CM 2

CRN 165252-70-0 CMF C7 H14 N4 O3

RN 886757-60-4 CAPLUS

CN Guanidine, N-methyl-N'-nitro-N''-[(tetrahydro-3-furanyl)methyl]-, mixt. with 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-dimethylpiperidine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-59-1 CMF C13 H19 N3 O S

$$Me-C = C-CH_2-O$$
 $N-S$
 Me
 Me
 Me

CM 2

CRN 165252-70-0 CMF C7 H14 N4 O3

RN 886757-62-6 CAPLUS

CN Guanidine, N-methyl-N'-nitro-N''-[(tetrahydro-3-furanyl)methyl]-, mixt. with 2,6-dimethyl-4-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]morpholine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-61-5 CMF C13 H19 N3 O2 S

CM 2

CRN 165252-70-0 CMF C7 H14 N4 O3

L12 ANSWER 10 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2006:489269 CAPLUS Full-text

DOCUMENT NUMBER:

144:462612

TITLE:

Pest control compositions containing cypermethrin and

pyrimidines or thiadiazoles

INVENTOR(S):

Shimokawatoko, Yasutaka; Yamada, Koji Sumitomo Chemical Co., Ltd., Japan

PATENT ASSIGNEE(S): SOURCE:

Jpn. Kokai Tokkyo Koho, 45 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent Japanese

LANGUAGE:

· 1

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.

KIND DATE

APPLICATION NO.

DATE

JP 2006131532

20060525 Α

JP 2004-321490 JP 2004-321490

20041105 20041105

PRIORITY APPLN. INFO.: OTHER SOURCE(S):

MARPAT 144:462612

GI

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Pest control compns. with excellent efficacy contain (1) pyrimidines (I, AΒ wherein R1 = H, alkyl; R2 = alkynyloxy; R3 = H, halo, alkyl; X1 = (un) substituted C4-7 polymethylene) or 1,2,4-thiadiazole derivs. such as 3-(2pentynyloxy)-5-(3-methylpiperidino)-1,2,4-thiadiazole and (2) (RS)- α -cyano-3phenoxybenzyl (1RS, 3RS; 1RS, 3SR) -3-(2, 2-dichlorovinyl) - 2, 2dimethylcyclopropanecarboxylate (cypermethrin). Thus, 4-(2-butynyloxy)-5fluoro-6-(3,3-dimethylpyrrolidin-1-yl)pyrimidine + cypermethrin at 12.5 + 1.5 ppm was more effective than the individual components at the same rates against Aphis gossypii on cucumber.

IT 886840-50-2 886840-51-3 886840-52-4 RL: AGR (Agricultural use); BSU (Biological study, unclassified); BIOL (Biological study); USES (Uses) (as pesticide)

RN 886840-50-2 CAPLUS

Cyclopropanecarboxylic acid, 3-(2,2-dichloroethenyl)-2,2-dimethyl-, CN cyano(3-phenoxyphenyl) methyl ester, mixt. with 3-methyl-1-[3-(2-methyl-1)]pentynyloxy)-1,2,4-thiadiazol-5-yl]piperidine (9CI) (CA INDEX NAME)

CM

850748-38-8 CRN CMF C13 H19 N3 O S

$$Et-C = C-CH_2-O$$
 $N-S$
 Me

CM

CRN 52315-07-8 CMF C22 H19 C12 N O3

RN 886840-51-3 CAPLUS

CN Cyclopropanecarboxylic acid, 3-(2,2-dichloroethenyl)-2,2-dimethyl-, cyano(3-phenoxyphenyl)methyl ester, mixt. with 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-dimethylpiperidine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-59-1 CMF C13 H19 N3 O S

$$Me-C = C-CH_2-O$$
 $N-S$
 Me
 Me
 Me

CM 2

CRN 52315-07-8 CMF C22 H19 C12 N O3

Me Me

RN 886840-52-4 CAPLUS

CN Cyclopropanecarboxylic acid, 3-(2,2-dichloroethenyl)-2,2-dimethyl-, cyano(3-phenoxyphenyl)methyl ester, mixt. with 2,6-dimethyl-4-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]morpholine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-61-5 CMF C13 H19 N3 O2 S

CRN 52315-07-8 CMF C22 H19 C12 N O3

Cl₂C CH CPh

L12 ANSWER 11 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2006:489268 CAPLUS Full-text

DOCUMENT NUMBER:

144:462611

TITLE:

Pest control compositions containing esfenvalerate and

pyrimidines or thiadiazoles

INVENTOR(S):

Shimokawatoko, Yasutaka; Yamada, Koji

PATENT ASSIGNEE(S):

Sumitomo Chemical Co., Ltd., Japan

SOURCE:

GΙ

Jpn. Kokai Tokkyo Koho, 45 pp.

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

CODEN: JKXXAF

FAMILY ACC. NUM. COUNT:

Japanes

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
 JP 2006131535	 A	20060525	JP 2004-321493	20041105
PRIORITY APPLN. INFO.: OTHER SOURCE(S):	маррат	144:462611	JP 2004-321493	20041105

AB Highly effective pest control compns. contain (1) pyrimidines (I, wherein R1 = H, alkyl; R2 = alkynyloxy; R3 = H, halo, alkyl; X1 = (un)substituted

polymethylene) or 1,2,4-thiadiazole derivs. such as 3-(2-pentynyloxy)-5-(3-methylpiperidino)-1,2,4-thiadiazole and (2) esfenvalerate as active components. Thus, <math>4-(2-butynyloxy)-5-fluoro-6-(3,3-dimethylpyrrolidin-1-yl)pyrimidine + Sumi-alpha emulsion at 12.5 + 1.5 ppm was more effective than the individual components at the same rates against Aphis gossypii on cucumber.

RN 886837-44-1 CAPLUS

CN Benzeneacetic acid, 4-chloro- α -(1-methylethyl)-, (S)-cyano(3-phenoxyphenyl)methyl ester, (α S)-, mixt. with 3-methyl-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]piperidine (9CI) (CA INDEX NAME)

CM 1

CRN 850748-38-8 CMF C13 H19 N3 O S

$$Et-C = C-CH_2-O$$
 $N-S$
 $N-S$

CM 2

CRN 66230-04-4 CMF C25 H22 C1 N O3

Absolute stereochemistry. Rotation (-).

RN 886837-46-3 CAPLUS

CN Benzeneacetic acid, 4-chloro- α -(1-methylethyl)-, (S)-cyano(3-phenoxyphenyl)methyl ester, (α S)-, mixt. with 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-dimethylpiperidine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-59-1 CMF C13 H19 N3 O S

$$Me-C = C-CH_2-O$$
 $N-S$
 Me
 Me

CRN 66230-04-4 CMF C25 H22 C1 N O3

Absolute stereochemistry. Rotation (-).

RN 886837-48-5 CAPLUS

CN Benzeneacetic acid, 4-chloro- α -(1-methylethyl)-, (S)-cyano(3-phenoxyphenyl)methyl ester, (α S)-, mixt. with 2,6-dimethyl-4-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]morpholine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-61-5 CMF C13 H19 N3 O2 S

CM 2

CRN 66230-04-4 CMF C25 H22 C1 N O3

Absolute stereochemistry. Rotation (-).

L12 ANSWER 12 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2006:489267 CAPLUS Full-text

DOCUMENT NUMBER:

144:462610

TITLE:

Pest control compositions containing spiromesifen and

pyrimidines or thiadiazoles

INVENTOR(S):

Shimokawatoko, Yasutaka; Yamada, Koji

PATENT ASSIGNEE(S):

Sumitomo Chemical Co., Ltd., Japan

SOURCE:

Jpn. Kokai Tokkyo Koho, 45 pp.

DOCUMENT TYPE:

Patent

LANGUAGE:

GI

RN

Japanese

CODEN: JKXXAF

FAMILY ACC. NUM. COUNT:

1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2006131531	Α	20060525	JP 2004-321489	20041105
PRIORITY APPLN. INFO.:			JP 2004-321489	20041105
OTHER SOURCE(S):	MARPAT	144:462610		

$$\begin{array}{c}
Me \\
N \\
\hline
N \\
\hline
N \\
O-CH2-C=C-CH3 \quad I
\end{array}$$

$$Et-C \equiv C-CH_2-O \xrightarrow{N-S} N \xrightarrow{Me} I$$

AB Compns. with excellent pest control effect contain (1) a pyrimidine derivative (e.g., I) or a 1,2,4-thiadiazole (e.g., II) and (2) 3-mesityl-2-oxo-1-oxaspiro[4.4]non-3-en-4-yl 3,3-dimethylbutylate (spiromesifen). Thus, I + spiromesifen at 0.15 + 6.3 ppm was more effective than the individual components at the same rates against Bemisia argentifolii on cabbage.

IT 886843-47-6 886843-48-7 886843-49-8
RL: AGR (Agricultural use); BSU (Biological study, unclassified); BIOL (Biological study); USES (Uses)

(as pesticide) 886843-47-6 CAPLUS

CN Butanoic acid, 3,3-dimethyl-, 2-oxo-3-(2,4,6-trimethylphenyl)-1-oxaspiro[4.4]non-3-en-4-yl ester, mixt. with 3-methyl-1-[3-(2-pentynyloxy)-

1,2,4-thiadiazol-5-yl]piperidine (9CI) (CA INDEX NAME)

CM 1

CRN 850748-38-8 CMF C13 H19 N3 O S

$$Et-C = C-CH_2-O$$

$$N-S$$
Me

CM 2

CRN 283594-90-1 CMF C23 H30 O4

RN 886843-48-7 CAPLUS

CN Butanoic acid, 3,3-dimethyl-, 2-oxo-3-(2,4,6-trimethylphenyl)-1-oxaspiro[4.4]non-3-en-4-yl ester, mixt. with 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-dimethylpiperidine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-59-1 CMF C13 H19 N3 O S

$$Me-C = C-CH_2-O$$
 N
 N
 Me
 N
 Me
 Me

CM 2

CRN 283594-90-1 CMF C23 H30 O4

RN 886843-49-8 CAPLUS

CN Butanoic acid, 3,3-dimethyl-, 2-oxo-3-(2,4,6-trimethylphenyl)-1-oxaspiro[4.4]non-3-en-4-yl ester, mixt. with 2,6-dimethyl-4-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]morpholine (9CI) (CA INDEX NAME)

CM 3

CRN 886757-61-5 CMF C13 H19 N3 O2 S

CM 2

CRN 283594-90-1 CMF C23 H30 O4

L12 ANSWER 13 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2006:489264 CAPLUS <u>Full-text</u>

DOCUMENT NUMBER:

144:462609

TITLE:

Pest control compositions containing acephate and

pyrimidines or thiadiazoles

INVENTOR(S):

Shimokawatoko, Yasutaka; Yamada, Koji Sumitomo Chemical Co., Ltd., Japan

PATENT ASSIGNEE(S):

Jpn. Kokai Tokkyo Koho, 44 pp.

SOURCE:

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT: 1 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2006131540	Α	20060525	JP 2004-321498	20041105
PRIORITY APPLN. INFO.:			JP 2004-321498	20041105
OTHER SOURCE(S):	MARPAT	144:462609		
GI				

AB Pest control compns. with excellent efficacy contain (1) a pyrimidine (I, wherein R1 = H, alkyl; R2 = alkynyloxy; R3 = H, halo, alkyl; X1 = (un)substituted C4-7 polymethylene) or a thiadiazole such as 3-(2-pentynyloxy)-5-(3-methylpiperidino)-1,2,4-thiadiazole and (2) O,S-di-Me acetylphosphoramidothioate (acephate). Thus, 4-(2-butynyloxy)-5-fluoro-6-(3,3-dimethylpyrrolidin-1-yl)pyrimidine + Ortran (500 + 500 ppm) gave almost 100% control of Aphis gossypii on cucumber.

IT 886973-61-1 886973-62-2 886973-63-3
RL: AGR (Agricultural use); BSU (Biological study, unclassified); BIOL (Biological study); USES (Uses)
(as pesticide)

RN 886973-61-1 CAPLUS

CN Phosphoramidothioic acid, acetyl-, O,S-dimethyl ester, mixt. with 3-methyl-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]piperidine (9CI) (CA INDEX NAME)

CM 1

CRN 850748-38-8 CMF C13 H19 N3 O S

$$Et-C = C-CH_2-O$$
 $N-S$
 $N-S$
 Me

CM 2

CRN 30560-19-1 CMF C4 H10 N O3 P S

RN 886973-62-2 CAPLUS

CN Phosphoramidothioic acid, acetyl-, O,S-dimethyl ester, mixt. with 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-dimethylpiperidine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-59-1 CMF C13 H19 N3 O S

$$Me-C = C-CH_2-O$$
 $N-S$
 Me
 Me
 Me

CM 2

CRN 30560-19-1 CMF C4 H10 N O3 P S

RN 886973-63-3 CAPLUS

CN Phosphoramidothioic acid, acetyl-, O,S-dimethyl ester, mixt. with 2,6-dimethyl-4-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]morpholine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-61-5 CMF C13 H19 N3 O2 S

CRN 30560-19-1 CMF C4 H10 N O3 P S

L12 ANSWER 14 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2006:489232 CAPLUS Full-text

DOCUMENT NUMBER: 144:462608

TITLE: Pest control compositions containing clothianidin and

pyrimidines or thiadiazoles

INVENTOR(S): Shimokawatoko, Yasutaka; Yamada, Koji

PATENT ASSIGNEE(S): Sumitomo Chemical Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 47 pp.

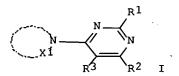
CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2006131534	Α	20060525	JP 2004-321492	20041105
PRIORITY APPLN. INFO.:			JP 2004-321492	20041105
OTHER SOURCE(S):	MARPAT	144:462608		
GI				



AB Pest control compns. with superior efficacy contain (1) a pyrimidine compound (I, wherein R1 = H, alkyl; R2 = alkynyloxy; R3 = H, halo, alkyl; X1 =

(un) substituted C4-7 polymethylene) or a 1,2,4-thiadiazole derivative such as 3-(2-pentynyloxy)-5-(3-methylpiperidino)-1,2,4-thiadiazole and (2) clothianidin as active components. Thus, 4-(2-butynyloxy)-5-fluoro-6-(3,3-dimethylpyrrolidin-1-yl) pyrimidine + clothianidin at 12.5 + 0.025 ppm was more effective than the individual components at the same rates against Aphis gossypii on cucumber.

IT 886758-19-6 886758-20-9 886758-21-0
RL: AGR (Agricultural use); BSU (Biological study, unclassified); BIOL (Biological study); USES (Uses)

(as pesticide) 886758-19-6 CAPLUS

CN Guanidine, N-[(2-chloro-5-thiazolyl)methyl]-N'-methyl-N''-nitro-, [C(E)]-, mixt. with 3-methyl-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]piperidine (9CI) (CA INDEX NAME)

CM 1

RN

CRN 850748-38-8 CMF C13 H19 N3 O S

$$Et-C = C-CH_2-O$$
N
N
N
Me

CM 2

CRN 210880-92-5 CMF C6 H8 C1 N5 O2 S

Double bond geometry as shown.

RN 886758-20-9 CAPLUS

CN Guanidine, N-[(2-chloro-5-thiazolyl)methyl]-N'-methyl-N''-nitro-, [C(E)]-, mixt. with 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-dimethylpiperidine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-59-1 CMF C13 H19 N3 O S

$$Me-C = C-CH_2-O$$
 N
 N
 Me
 Me

CM 2

CRN 210880-92-5 CMF C6 H8 Cl N5 O2 S

Double bond geometry as shown.

RN 886758-21-0 CAPLUS

CN Guanidine, N-[(2-chloro-5-thiazolyl)methyl]-N'-methyl-N''-nitro-, [C(E)]-, mixt. with 2,6-dimethyl-4-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]morpholine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-61-5 CMF C13 H19 N3 O2 S

$$Et-C = C-CH_2-O \xrightarrow{N} \xrightarrow{Me} O$$

CM 2

CRN 210880-92-5 CMF C6 H8 Cl N5 O2 S

Double bond geometry as shown.

L12 ANSWER 15 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2006:489228 CAPLUS Full-text

DOCUMENT NUMBER:

144:462607

TITLE:

Pesticides containing buprofezin and a pyrimidine or

thiadiazole derivative

INVENTOR(S):

Shimokawatoko, Yasutaka; Yamada, Koji Sumitomo Chemical Co., Ltd., Japan

PATENT ASSIGNEE(S):

Jpn. Kokai Tokkyo Koho, 45 pp.

SOURCE:

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2006131527 PRIORITY APPLN. INFO.:	A .	20060525	JP 2004-321485 JP 2004-321485	20041105 20041105
OTHER SOURCE(S):	MARPAT	144:462607		

$$\begin{array}{c}
Me \\
N \\
N \\
N
\end{array}$$

$$\begin{array}{c}
N \\
N \\
O-CH2-C \equiv C-CH3
\end{array}$$
I

$$Et-C \equiv C-CH_2-O \nearrow N \longrightarrow N \longrightarrow Me$$

- AB Compns. with excellent pest control effect comprise (1) a pyrimidine compound (e.g., I) or a 1,2,4-thiadiazole compound (e.g., II) and (2) buprofezin. Thus, I + buprofezin at 0.15 + 3.1 ppm showed greater control of Bemisia argentifolii on cabbage than the individual components at the same resp.
- RN 886842-21-3 CAPLUS
- CN 4H-1,3,5-Thiadiazin-4-one, 2-[(1,1-dimethylethyl)imino]tetrahydro-3-(1-methylethyl)-5-phenyl-, mixt. with 3-methyl-1-[3-(2-pentynyloxy)-1,2,4-

thiadiazol-5-yl]piperidine (9CI) (CA INDEX NAME)

CM 1

CRN 850748-38-8 CMF C13 H19 N3 O S

$$Et-C = C-CH_2-O$$
 $N-S$
 $N-S$
 Me

CM 2

CRN 69327-76-0 CMF C16 H23 N3 O S

RN 886842-22-4 CAPLUS

CN 4H-1,3,5-Thiadiazin-4-one, 2-[(1,1-dimethylethyl)imino]tetrahydro-3-(1-methylethyl)-5-phenyl-, mixt. with 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-dimethylpiperidine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-59-1 CMF C13 H19 N3 O S

$$Me-C = C-CH_2-O$$
 $N-S$
 Me
 Me
 Me

CM 2

CRN 69327-76-0 CMF C16 H23 N3 O S

RN886842-23-5 CAPLUS

CN 4H-1,3,5-Thiadiazin-4-one, 2-[(1,1-dimethylethyl)imino]tetrahydro-3-(1methylethyl)-5-phenyl-, mixt. with 2,6-dimethyl-4-[3-(2-pentynyloxy)-1,2,4thiadiazol-5-yl]morpholine (9CI) (CA INDEX NAME)

CM

CRN 886757-61-5 CMF C13 H19 N3 O2 S

CM 2

CRN 69327-76-0 CMF C16 H23 N3 O S

L12 ANSWER 16 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN ACCESSION NUMBER: 2006:489226 CAPLUS Full-text

DOCUMENT NUMBER: 144:462606

TITLE: Pest control compositions containing pymetrozine and

pyrimidines or thiadiazoles

INVENTOR(S): Shimokawatoko, Yasutaka; Yamada, Koji

PATENT ASSIGNEE(S): Sumitomo Chemical Co., Ltd., Japan

Jpn. Kokai Tokkyo Koho, 45 pp. SOURCE:

CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese FAMILY ACC. NUM. COUNT: 1 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
		 -		
JP 2006131529	Α	20060525	JP 2004-321487	20041105
PRIORITY APPLN. INFO.:			JP 2004-321487	20041105
OTHER SOURCE(S):	MARPAT	144:462606		
GI				

AB Pest control compns. with excellent effect contain (1) a pyrimidine compound (e.g., I) or a 1,2,4-thiadiazole derivative (e.g., II) and (2) (E)-4,5-dihydro-6-methyl-4-[(3-pyridylmethylene)amino]-1,2,4-triazin-3(2H)- one (pymetrozine) as active components. Thus, I + pymetrozine at 12.5 + 0.1 ppm gave better control of Aphis gossypii on cucumber than did the individual components at the same resp. rates.

RN 886842-10-0 CAPLUS

CN 1,2,4-Triazin-3(2H)-one, 4,5-dihydro-6-methyl-4-[(E)-(3-pyridinylmethylene)amino]-, mixt. with 3-methyl-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]piperidine (9CI) (CA INDEX NAME)

CM 1

CRN 850748-38-8 CMF C13 H19 N3 O S

$$Et-C \equiv C-CH_2-O$$
 $N-S$
 Me

CM 2

CRN 123312-89-0 CMF C10 H11 N5 O Double bond geometry as shown.

RN 886842-11-1 CAPLUS

CN 1,2,4-Triazin-3(2H)-one, 4,5-dihydro-6-methyl-4-[(E)-(3-pyridinylmethylene)amino]-, mixt. with 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-dimethylpiperidine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-59-1 CMF C13 H19 N3 O S

$$Me-C = C-CH_2-O$$
 $N-S$
 Me
 Me
 Me

CM 2

CRN 123312-89-0 CMF C10 H11 N5 O

Double bond geometry as shown.

RN 886842-12-2 CAPLUS

CN 1,2,4-Triazin-3(2H)-one, 4,5-dihydro-6-methyl-4-[(E)-(3-pyridinylmethylene)amino]-, mixt. with 2,6-dimethyl-4-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]morpholine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-61-5

CMF C13 H19 N3 O2 S

CM 2

CRN 123312-89-0 CMF C10 H11 N5 O

Double bond geometry as shown.

L12 ANSWER 17 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2006:489222 CAPLUS Full-text

DOCUMENT NUMBER:

144:462605

TITLE:

Pesticides containing flonicamid and pyrimidine or

thiadiazole derivative

INVENTOR(S):

Shimokawatoko, Yasutaka; Yamada, Koji

PATENT ASSIGNEE(S):

Sumitomo Chemical Co., Ltd., Japan

SOURCE:

Jpn. Kokai Tokkyo Koho, 45 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2006131526	Α	20060525	JP 2004-321484	20041105
PRIORITY APPLN. INFO.:			JP 2004-321484	20041105
OTHER SOURCE(S):	MARPAT	144:462605		
GI				

$$N \longrightarrow \mathbb{R}^{1}$$
 \mathbb{R}^{2}
 \mathbb{R}^{2}
 \mathbb{R}^{2}

Compns. with excellent pest control effect contain (1) a pyrimidine derivative (I, where R1 and R3 = H, etc.; R2 = C3-7 alkynyloxy; X1 = C4-7 polymethylene, etc.) or a 1,2,4-thiadiazole derivative (e.g., 3-(2-pentynyloxy)-5-(3-methylpiperidino)-1,2,4-thiadiazole) and (2) flonicamid as active components. Thus, 4-(2-butynyloxy)-5-fluoro-6-(3,3-dimethylpyrrolidin-1-yl)pyrimidine + flonicamid at 6.3 + 0.8 ppm was more effective than the individual components at the same rates against Aphis gossypii on cucumber.

IT 886843-94-3 886843-95-4 886843-96-5
RL: AGR (Agricultural use); BSU (Biological study, unclassified); BIOL (Biological study); USES (Uses)
(as pesticide)

RN 886843-94-3 CAPLUS

CN 3-Pyridinecarboxamide, N-(cyanomethyl)-4-(trifluoromethyl)-, mixt. with 3-methyl-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]piperidine (9CI) (CA INDEX NAME)

CM 1

CRN 850748-38-8 CMF C13 H19 N3 O S

$$Et-C = C-CH_2-O$$
 $N-S$
 $N-S$

CM 2

CRN 158062-67-0 CMF C9 H6 F3 N3 O

RN 886843-95-4 CAPLUS

CN 3-Pyridinecarboxamide, N-(cyanomethyl)-4-(trifluoromethyl)-, mixt. with 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-dimethylpiperidine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-59-1 CMF C13 H19 N3 O S

$$Me-C = C-CH_2-O$$
 $N-S$
 Me
 Me

CM 2

CRN 158062-67-0 CMF C9 H6 F3 N3 O

$$NC-CH_2-NH-C$$
 F_3C

RN 886843-96-5 CAPLUS

CN 3-Pyridinecarboxamide, N-(cyanomethyl)-4-(trifluoromethyl)-, mixt. with 2,6-dimethyl-4-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]morpholine (9CI) (CA INDEX NAME)

CM 1

CRN 886757-61-5 CMF C13 H19 N3 O2 S

$$Et-C = C-CH_2-O \xrightarrow{N} \xrightarrow{Me} O$$

CM 2

CRN 158062-67-0 CMF C9 H6 F3 N3 O

L12 ANSWER 18 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2006:34718 CAPLUS <u>Full-text</u>

DOCUMENT NUMBER: 144:108364

TITLE: Preparation of 1-(1-oxo-2-propynyl)piperazines as

mGLuR5 receptor modulators for the treatment of pain

INVENTOR(S): Kuehnert, Sven; Oberboersch, Stefan; Haurand, Michael;

Jostock, Ruth; Schiene, Klaus

PATENT ASSIGNEE(S): Gruenthal GmbH, Germany

SOURCE: PCT Int. Appl., 208 pp.

CODEN: PIXXD2
DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PA	TENT	NO.			KIN	D	DATE								Di	ATE	
WO	2006	0029	81		A1	_	2006	0112	1		: 005-:				2	0050	705
	W:	ΑE,	AG,	AL,	AM,	ΑT,	ΑU,	ΑZ,	BA,	BB,	BG,	BR,	BW,	BY,	ΒZ,	CA,	CH,
							DK,										
		GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	ΚE,	KG,	KM,	ΚP,	KR,	ΚZ,	LC,
		LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NA,	NG,
		NI,	NO,	NZ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,
		SM,	SY,	TJ,	TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VC,	VN,	YU,	ZA,
		ZM,	zw														
	RW:	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	ΙE,
		IS,	IT,	LT,	LU,	LV,	MC,	NL,	PL,	PT,	RO,	SE,	SI,	SK,	TR,	BF,	ВJ,
		CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,	ΝE,	SN,	TD,	TG,	BW,	GH,
		GM,	ΚE,	LS,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	ΑZ,	BY,
		KG,	ΚZ,	MD,	RU,	ТJ,	TM										
DE	1020	0403	2567		A 1		2006	0302		DE 2	004-	1020	0403	2567	2	0040	705
. CA	. 2572	685			A1		2006	0112		CA 2	005-	2572	685		2	0050	705
EP	1765	816			A1		2007	0328		EP 2	005-	7565	39		2	0050	705
	R:	ΑT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	ΙE,
		IS,	IT,	LI,	LT,	LU,	LV,	MC,	NL,	PL,	PΤ,	RO,	SE,	SI,	SK,	TR	
US	2007	1120	11		A1		2007	0517		US 2	007-	6491	56		2	0070	104
US	7300	939			В2	•	2007	1127									
PRIORIT	Y APP	LN.	INFO	.:						DE 2	004-	1020	0403	2567.	A 2	0040	705
									1	WO 2	005-	EP72	48	1	W 2	0050	705
OTHER S	OURCE	(S):			CAS	REAC	Т 14	4:10	8364	; MA	RPAT	144	:108	364			

46

$$N - CO - C \equiv C - R4$$
 $N - CO - C \equiv C - CH_2Ph$

III

AB Title compds. I [Z = (R3)n; X = N, CR2; R1, R2 = H, halo, NO2, etc.; R3 = halo, NO2, CN, etc.; R4 = H, halo, NO2, etc.] and their pharmaceutically acceptable salts were prepared For example, coupling of 4-phenyl-2-butynoic acid and 1-thiazol-2-ylpiperazine afforded claimed propynylpiperazine II in79% yield. In mGLuR5 receptor binding assays, propynylpiperazine III exhibited an IC50 value of 100 nM.

IT 873073-78-0P 873073-80-4P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of 1-(1-oxo-2-propynyl)) piperazines as mGLuR5 receptor modulators for the treatment of pain)

RN 873073-78-0 CAPLUS

CN Piperazine, 1-[3-(methylthio)-1,2,4-thiadiazol-5-yl]-4-(1-oxo-3-phenyl-2-propynyl)- (9CI) (CA INDEX NAME)

$$Mes \xrightarrow{N} S \xrightarrow{N} C = C - Ph$$

RN 873073-80-4 CAPLUS

CN Piperazine, 1-[3-(methylsulfonyl)-1,2,4-thiadiazol-5-yl]-4-(1-oxo-3-phenyl-2-propynyl)- (9CI) (CA INDEX NAME)

$$Me = \bigcup_{N=1}^{\infty} \bigvee_{N=1}^{N} \bigvee_{N=1}^{\infty} \bigcup_{N=1}^{\infty} C = Ph$$

RN 873075-56-0 CAPLUS

CN Piperazine, 1-[3-(methylsulfonyl)-1,2,4-thiadiazol-5-yl]- (CA INDEX NAME)

$$Me-\bigcup_{N=1}^{O} \bigvee_{N=1}^{N} \bigvee$$

REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L12 ANSWER 19 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN ACCESSION NUMBER: 2005:371234 CAPLUS Full-text

DOCUMENT NUMBER:

142:430280

TITLE:

Preparation of 1,2,4-thiadiazole compounds

as pests controlling agents

INVENTOR(S):

Ihara, Hideki; Takaoka, Daisuke;

Mizuno, Hajime

PATENT ASSIGNEE(S):

Sumitomo Chemical Company, Limited, Japan

SOURCE: PCT Int. Appl., 55 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PA'	TENT	NO.			KIN	D	DATE			APPL	ICAT	ION 1	NO.		D.	ATE	
	2005 2005						2005 2007		,	WO 2	004-	JP14	540		2	0040	927
	W:	ΑE,	AG,	AL,	AM,	ΑT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BW,	BY,	BZ,	CA,	CH,
							DE,										
							ID,										
							MA,										
							PT,										
		TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	zw	•
	RW:	BW,	GH,	GM,	KE,	LS,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,
							RU,										
							GR,										
							CF,										
•							EP,								•	•	•
AU	2004	2820	18		A1		2005	0428		AU 2	004-	2820	18		2	0040	927
BR	2004	0153	64 .		Α		2006	1212		BR 2	004-	1536	4		2	0040	927
	1765															0040	927
	R:	ΑT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	IE,
		IT,	LI,	LU,	MC,	NL,	PL,	PT,	RO,	SE,	SI,	SK,	TR				
JP	2005	1391	71		Α		2005	0602		JP 2	004-	2973	25		2	0041	012
US	2007	0047	22		A 1		2007	0104	,	US 2	006-	5679	84		2	0060	210
MX	2006	PA04	118		Α		2006	0705]	MX 2	006-	PA41	18		2	0060	411
IN	2006	CN01	272		Α		2007	0629		IN 2	006-	CN12	72		2	0060	413
PRIORIT	Y APP	LN.	INFO	.:						JP 2	003-	3547	58		A 2	0031	015
									1	WO 2	004-	JP14	540	1	w 2	0040	927
OTHER SO	OURCE	(S):			CAS	REAC	т 14	2:43	0280	; MA	RPAT	142	:430	280			

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$$R^{1}-O$$
 $N-S$
 $N-S$
 $N-S$
 $N-S$

AB Title compds. I [R1 = alkynyl; X = (un)substituted straight alkylene, (un)substituted straight alkenylene, (un)substituted ethylene-oxyethylene, etc.] were prepared For example, aromatic nucleophilic substitution of 5-chloro-3-methylthio-1,2,4-thiadiazole with pyrrolidine followed by oxidation using 3-chloroperbenzoic acid and treatment with 2-butyn-1-ol afforded 3-(2-butynyloxy)-5-(pyrrolidin-1-yl)-1,2,4-thiadiazole. In pest controlling test against aphis gossypii, compound II had the control value of ≥90%. Compds. I are claimed useful as pests controlling agents. Formulations are given.

TT 850748-31-1P 850748-32-2P 850748-33-3P 850748-34-4P 850748-35-5P 850748-36-6P 850748-37-7P 850748-38-8P 850748-42-4P 850748-43-5P 850748-44-6P 850748-45-7P 850748-46-8P 850748-47-9P 850748-49-1P 850748-50-4P 850748-51-5P 850748-52-6P 850748-53-7P 850748-55-9P 850748-55-1P

850748-58-2P RL: AGR (Agricultural use); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of 1,2,4-thiadiazole compds. as pests controlling agents)

RN 850748-31-1 CAPLUS

CN 1,2,4-Thiadiazole, 3-(2-butynyloxy)-5-(1-pyrrolidinyl)- (9CI) (CA INDEX NAME)

$$Me-C = C-CH_2-O$$
 $N-S$

RN 850748-32-2 CAPLUS
CN 1,2,4-Thiadiazole, 3-(2-pentynyloxy)-5-(1-pyrrolidinyl)- (9CI) (CA INDEX NAME)

$$Et-C = C-CH_2-O$$
N
N
N
N

RN 850748-33-3 CAPLUS

CN Piperidine, 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]- (9CI) (CA INDEX NAME)

$$Me-C = C-CH_2-O$$
 $N-S$

RN 850748-34-4 CAPLUS

CN Piperidine, 1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]- (9CI) (CA INDEX NAME)

$$Et-C = C-CH_2-O NS N$$

RN 850748-35-5 CAPLUS

CN Piperidine, 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-2-methyl- (9CI) (CA INDEX NAME)

RN 850748-36-6 CAPLUS

CN Piperidine, 2-methyl-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]- (9CI) (CA INDEX NAME)

$$Et-C = C-CH_2-O \xrightarrow{N} \xrightarrow{Me} N$$

RN 850748-37-7 CAPLUS

CN Piperidine, 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3-methyl- (9CI)

(CA INDEX NAME)

$$Me-C = C-CH_2-O$$
 $N-S$
 $N-S$

RN 850748-38-8 CAPLUS

CN Piperidine, 3-methyl-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]- (9CI) (CA INDEX NAME)

$$Et-C = C-CH_2-O N N N$$

RN 850748-39-9 CAPLUS

CN Morpholine, 4-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]- (9CI) (CA INDEX NAME)

$$Me-C = C-CH_2-O$$
 $N-S$

RN 850748-40-2 CAPLUS

CN Morpholine, 4-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]- (9CI) (CA INDEX NAME)

RN 850748-41-3 CAPLUS

CN Piperidine, 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-4-methyl- (9CI) (CA INDEX NAME)

$$Me-C = C-CH_2-O \xrightarrow{N} N$$

RN 850748-42-4 CAPLUS

CN Piperidine, 4-methyl-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]- (9CI) (CA INDEX NAME)

$$Et-C = C-CH_2-O$$

$$N-S$$

$$N-S$$

RN 850748-43-5 CAPLUS

CN 1H-Azepine, 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]hexahydro- (9CI) (CA INDEX NAME)

$$Me-C = C-CH_2-O$$
 $N-S$
 $N-S$

RN 850748-44-6 CAPLUS

CN 1H-Azepine, hexahydro-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]- (9CI) (CA INDEX NAME)

RN 850748-45-7 CAPLUS

CN Piperidine, 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-dimethyl-, (3R,5S)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.

RN 850748-46-8 CAPLUS

CN Piperidine, 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-dimethyl-, (3R,5R)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.

RN 850748-47-9 CAPLUS

CN Piperidine, 3,5-dimethyl-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]-, (3R,5S)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.

RN 850748-48-0 CAPLUS

CN Piperidine, 3,5-dimethyl-1-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]-, (3R,5R)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.

RN 850748-49-1 CAPLUS

CN Morpholine, 4-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-2,6-dimethyl-, (2R,6S)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.

RN 850748-50-4 CAPLUS

CN Morpholine, 4-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-2,6-dimethyl-, (2R,6R)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.

RN 850748-51-5 CAPLUS

CN Morpholine, 2,6-dimethyl-4-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]-, (2R,6S)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.

RN 850748-52-6 CAPLUS

CN Morpholine, 2,6-dimethyl-4-[3-(2-pentynyloxy)-1,2,4-thiadiazol-5-yl]-, (2R,6R)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.

RN 850748-53-7 CAPLUS

CN Piperidine, 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-diethyl-, (3R,5S)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.

RN 850748-54-8 CAPLUS

CN Piperidine, 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3,5-diethyl-, (3R,5R)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.

$$Me _C = C$$

$$N$$

$$N$$

$$R$$

$$Et$$

$$N$$

$$Et$$

RN 850748-55-9 CAPLUS

CN Piperidine, 3,5-dimethyl-1-[3-(2-propynyloxy)-1,2,4-thiadiazol-5-yl]-(9CI) (CA INDEX NAME)

$$HC = C - CH_2 - O$$
 $N - S$
 Me
 Me
 Me

RN 850748-56-0 CAPLUS

CN Piperidine, 3,5-dimethyl-1-[3-[(1-methyl-2-butynyl)oxy]-1,2,4-thiadiazol-5-yl]- (9CI) (CA INDEX NAME)

$$Me-C = C-CH-O$$
 $N-S$
 Me
 Me
 Me
 Me

RN 850748-57-1 CAPLUS

CN Piperidine, 1-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]-3-(trifluoromethyl)-(9CI) (CA INDEX NAME)

$$Me-C = C-CH_2-O$$
 $N-S$
 $N-S$
 $N-S$

RN 850748-58-2 CAPLUS

CN Thiomorpholine, 4-[3-(2-butynyloxy)-1,2,4-thiadiazol-5-yl]- (9CI) (CA INDEX NAME)

IT 850748-59-3P 850748-60-6P 850748-61-7P

850748-62-8P 850748-63-9P 850748-64-0P

850748-65-1P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of 1,2,4-thiadiazole compds. as pests controlling

agents)

RN 850748-59-3 CAPLUS

CN Piperidine, 3,5-dimethyl-1-[3-(methylthio)-1,2,4-thiadiazol-5-yl]- (CA INDEX NAME)

RN 850748-60-6 CAPLUS

CN Piperidine, 3,5-dimethyl-1-[3-(methylsulfonyl)-1,2,4-thiadiazol-5-yl]-(CA INDEX NAME)

$$Me = \bigcup_{N=-S}^{O} \bigvee_{N=-S}^{Me} \bigvee_{Me}^{Me}$$

RN 850748-61-7 CAPLUS

CN Piperidine, 3,5-dimethyl-1-[3-(methylsulfinyl)-1,2,4-thiadiazol-5-yl]-(CA INDEX NAME)

RN 850748-62-8 CAPLUS

CN Piperidine, 1-[3-(methylthio)-1,2,4-thiadiazol-5-yl]-3-(trifluoromethyl)-(CA INDEX NAME)

RN 850748-63-9 CAPLUS

CN Piperidine, 1-[3-(methylsulfonyl)-1,2,4-thiadiazol-5-yl]-3-(trifluoromethyl)- (CA INDEX NAME)

$$Me-\bigcup_{N=1}^{O} \bigvee_{N=1}^{N} \bigvee_{N=1}^{N} \bigvee_{N=1}^{N} CF3$$

RN 850748-64-0 CAPLUS

CN Thiomorpholine, 4-[3-(methylsulfonyl)-1,2,4-thiadiazol-5-yl]- (CA INDEX NAME)

$$Me-\bigcup_{N-S}^{\circ} N \longrightarrow N$$

RN 850748-65-1 CAPLUS

CN Thiomorpholine, 4-[3-(methylsulfinyl)-1,2,4-thiadiazol-5-yl]- (CA INDEX NAME)

$$Me-\overset{\circ}{\underset{N-S}{\parallel}} \overset{\circ}{\underset{N-S}{\parallel}} \overset{\circ}{\underset{N}{\longrightarrow}} \overset{N}{\longrightarrow} \overset{\sim}{\underset{N}{\longrightarrow}} \overset{\sim}{\underset{N}{\longrightarrow}} \overset{\sim}{\underset{N}{\longrightarrow}} \overset{\sim}{\underset{N}{\longrightarrow}} \overset{N}{\longrightarrow} \overset{\sim}{\underset{N}{\longrightarrow}} \overset{\sim}{\underset{N}{\longrightarrow}} \overset{\sim}{\underset{N}{\longrightarrow}} \overset{\sim}{\underset{N}{\longrightarrow}} \overset{N}{\longrightarrow} \overset{\sim}{\underset{N}{\longrightarrow}} \overset{\sim}{\underset{N}{\longrightarrow}} \overset{\sim}{\underset{N}{\longrightarrow}} \overset{\sim}{\underset{N}{\longrightarrow}} \overset{N}{\longrightarrow}} \overset{\sim}{\underset{N}{\longrightarrow}} \overset{\sim}{\underset{N}{\longrightarrow}} \overset{\sim}{\underset{N}{\longrightarrow}} \overset{\sim}{\underset{N}{\longrightarrow}} \overset{$$

L12 ANSWER 20 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN ACCESSION NUMBER: 2004:182866 CAPLUS Full-text

DOCUMENT NUMBER: 140:236096

TITLE: Preparation of proline derivatives as antibacterial

agents

INVENTOR(S): Fujita, Masahiro; Sakamoto, Masato; Horiuchi,

Nobuhiko; Yamamoto, Takayoshi; Tomita, Kyoji; Mizuno, Kazuhiro; Niga, Toshiyuki; Ito, Hideaki; Kashimoto,

Shigeki

PATENT ASSIGNEE(S): Dainippon Pharmaceutical Co., Ltd., Japan

SOURCE: PCT Int. Appl., 122 pp.

CODEN: PIXXD2

DOCUMENT TYPE: LANGUAGE:

Patent Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PA'	TENT 1	NO.			KIN	D :	DATE			APP)	LICAT	ION 1	NO.			ATE	
WO	2004	0184	53		A1	_	2004	0304	,	WO 2	2003-	JP10	 548				
	W:	ΑE,	AG,	AL,	AM,	AT,	AU,	AZ,	BA,	BB,	, BG,	BR,	BY,	BZ,	CA,	CH,	CN,
		co,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	, EE,	ES,	FI,	GB,	GD,	GE,	GH,
		GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	, KG,	KR,	ΚZ,	LC,	LK,	LR,	LS,
		LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	, MX,	MZ,	NI,	NO,	NZ,	OM,	PG,
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		TT,	TZ,	UA,	UG,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	ZW				
	RW:	GH,	GM,	ΚE,	LS,	MW,	MZ,	SD,	SL,	SZ	, TZ,	UG,	ZM,	ZW,	AM,	ΑZ,	BY,
		KG,	ΚZ,	MD,	RU,	ТJ,	TM,	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,
		FI,	FR,	GB,	GR,	HU,	ΙE,	IT,	LU,	MC,	, NL,	PT,	RO,	SE,	SI,	SK,	TR,
		BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	, GW,	ML,	MR,	NE,	SN,	TD,	TG
JP	2006	0521	38		Α	:	2006	0223		JP 2	2002-	2427	95		-2	0020	823
JP	2006	0521	39		Α		2006	0223		JP 2	2002-	3392	00		2	0021	122
JP	2006	0521	40		Α		2006	0223		JP 2	2003-	2701	0		2	0030	204
AU	2003	2576	37		A 1		2004	0311	_	AU 2	2003-	2576	37		2	0030	821
PRIORIT	Y APP	LN.	INFO	.:						JP 2	2002÷	2427	95	i	A 2	0020	823
										JP 2	2002-	3392	00	i	A 2	0021	122
										JP 2	2003-	2701	0	1	A 2	0030	204
									1	WO 2	2003-	JP10	548	1	w 2	0030	821

OTHER SOURCE(S):

MARPAT 140:236096

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$$\begin{array}{c|c} & & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & \\ & & \\$$

AΒ Proline derivs. represented by the general formula (I) or salts thereof [wherein A = a group derived from a 5- or 6-membered heterocycle which may be fused with an optionally halogenated benzene ring; p1, p2, p3 = 0, 1; R1, R2, R3 = H, lower alkoxy, lower alkylthio, halo, HO, (un)protected or (un) substituted NH2 or CONH2, hydroxy-lower alkylamino, CO2H, lower alkoxycarbonyl, lower alkylcarbonyloxy, (un)substituted lower alkylsulfonyloxy, cyano; when p1 = p2 = 1, CR1R2 = CO; or when p1 = p2 = p3 =1, R1 = R2 = H and R3 = a 5- or 6-membered saturated or unsatd. cyclic group; T = a single bond, CH2, CO; R4, R5 = H, lower alkyl; or CR4R5 = CO; n, m =1,2; R6, R7 = H, OH, halogeno, lower alkyl, Ph, lower alkoxy, phenyl-lower alkyl, (un)protected NH2; R6 and R7 together form a saturated cyclic group; X = CH2, CH, S, O; Y = H, an amino-protecting group, or a group represented by the general formula R9ON(CHO)CH2CH(R8)CO; wherein R8 = alkyl, cycloalkyl-lower alkyl; R9 = H, a hydroxyl-protecting group, etc.] are prepared These compds. are useful as antibacterial drugs against multidrug-resistant bacteria. Thus, (2R) -3-cyclopentyl-2-[N-(2,4-dimethoxybenzyloxy)-Nformylamino]methyl]propionic acid was condensed with (2S)-2-[[4-(2pyrimidinyl)-1-piperazinyl]carbonyl]pyrrolidine hydrochloride using 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide hydrochloride, 1-hydroxybenzotriazole, and Et3N in CH2Cl2 at room temperature for 18 h to give 68% (2S)-1-[(2R)-3cyclopentyl-2-[[N-(2,4- dimethoxybenzyloxy)-N-formylamino]methyl]propionyl]-2-[[4-(2-pyrimidinyl)- 1-piperazinyl]carbonyl]pyrrolidine which was treated with 3% CF3CO2H in CH2Cl2 at room temperature for 17 h and then with saturated aqueous NaHCO3 under ice-cooling to give 77% (2S)-1-[(2R)-3-cyclopentyl-2-[(Nformyl-N- hydroxyamino)methyl]propionyl]-2-[[4-(2-pyrimidinyl)-1piperazinyl]carbonyl]pyrrolidine (II). II showed min. inhibitory concentration of 0.25, 0.125, 0.03, 0.25, 0.5, 0.125, 1, 0.5, and 0.125 $\mu g/mL$ against Staphylococcus aureus Smith, S. aureus KTO150 (MRSA), S. epidermidis ATCC12228, Streptococcus pneumoniae ATCC49619, S. pneumoniae KT2524 (PRSP), S. pneumoniae KB2534 (PRSP), S. pyrogenes ATCC12344, Enterococcus faecium ATCC19434, and Moraxella (B.) catarrhalis K1209, resp.

IT 668483-33-8P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of proline derivs. as antibacterial agents against multidrug-resistant bacteria)

RN 668483-33-8 CAPLUS

CN Piperazine, $1-[(2R)-N-formyl-N-hydroxy-2-pentyl-\beta-alanyl-L-prolyl]-4-$

[3-(methylthio)-1,2,4-thiadiazol-5-yl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

IT 668484-30-8P 668484-89-7P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of proline derivs. as antibacterial agents against multidrug-resistant bacteria)

RN 668484-30-8 CAPLUS

CN 1-Piperazinecarboxylic acid, 4-[3-(methylthio)-1,2,4-thiadiazol-5-yl]-, 1,1-dimethylethyl ester (CA INDEX NAME)

RN 668484-89-7 CAPLUS

CN 1-Pyrrolidinecarboxylic acid, 2-[[4-[3-(methylthio)-1,2,4-thiadiazol-5-yl]-1-piperazinyl]carbonyl]-, 1,1-dimethylethyl ester, (2S)- (CA INDEX NAME)

Absolute stereochemistry.

REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L12 ANSWER 21 OF 21 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

1972:113135 CAPLUS Full-text

DOCUMENT NUMBER:

76:113135

ORIGINAL REFERENCE NO.:

76:18269a,18272a

TITLE:

Organic herbicides. V. Derivatives of 3-mercapto-5-amino-1,2,4-thiadiazole Zbirovsky, M.: Myska, J.: Stanek, J.

AUTHOR(S):

Zbirovsky, M.; Myska, J.; Stanek, J. Vysk. Sk. Chem. Technol., Prague, Czech.

CORPORATE SOURCE: SOURCE:

Collection of Czechoslovak Chemical Communications

(1971), 36(12), 4087-91

CODEN: CCCCAK; ISSN: 0010-0765

DOCUMENT TYPE:

Journal

LANGUAGE:

English

GI For diagram(s), see printed CA Issue.

AB -MeS, 3-EtS, 3-allylthio, and 3-PhCH2S, derivs. of 5-chloro-1,2,4- thiadiazole were refluxed in MeOH with MeNH2, EtNH2, iso-PrNH2, BuNH2, CH2:CHCH2NH2, PhNH2, piperidine, or morpholine to give 27 S3,N5-disubstituted-3-mercapto-5-amino-1,2,4-thiadiazoles (I) the herbicidal activity of which was too low for practical use.

IT 35746-49-7P 35746-55-5P 35746-61-3P

RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of)

RN 35746-49-7 CAPLUS

CN Morpholine, 4-[3-(methylthio)-1,2,4-thiadiazol-5-yl]- (CA INDEX NAME)

RN 35746-55-5 CAPLUS

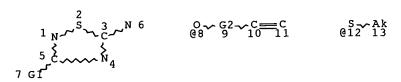
CN Morpholine, 4-[3-(ethylthio)-1,2,4-thiadiazol-5-yl]- (CA INDEX NAME)

RN 35746-61-3 CAPLUS

CN Morpholine, 4-[3-(2-propenylthio)-1,2,4-thiadiazol-5-yl]- (9CI) (CA INDEX NAME)

$$H_2C = CH - CH_2 - S$$
 $N - S$
 $N - S$

=> d que 113 STR



VAR G1=8/12REP G2 = (0-10) C NODE ATTRIBUTES:

NSPEC IS R ΑT CONNECT IS E1 RC AT 13 DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 13

STEREO ATTRIBUTES: NONE

010		
L3	98	SEA FILE=REGISTRY SSS FUL L1
L4	21	SEA FILE=CAPLUS ABB=ON PLU=ON L3
L5	242	SEA FILE=CAPLUS ABB=ON PLU=ON ("IHARA H"/AU OR "IHARA
		HIDEAKI"/AU OR "IHARA HIDEKI"/AU)
L6	51	SEA FILE=CAPLUS ABB=ON PLU=ON ("TAKAOKA D"/AU OR "TAKAOKA
		DAISUKE"/AU)
L7	186	SEA FILE=CAPLUS ABB=ON PLU=ON ("MIZUNO H"/AU OR "MIZUNO
		HAJIME"/AU)
L8	474	SEA FILE=CAPLUS ABB=ON PLU=ON (L5 OR L6 OR L7)
L9	6	SEA FILE=CAPLUS ABB=ON. PLU=ON L8 AND THIA? AND ?AZOL?
L10	1	SEA FILE=CAPLUS ABB=ON PLU=ON L9 AND L4
т.1.3	5	SEA FILE=CAPLUS ABB=ON PLU=ON L9 NOT L10

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L13 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2007 ACS on STN ACCESSION NUMBER: 2004:453198 CAPLUS Full-text

DOCUMENT NUMBER: 141:23538

1,2,4-Thiadiazole compounds, their TITLE:

preparation, their use as pesticides, and

arthropod-controlling compositions containing them

Ihara, Hideki; Takaoka, Daisuke INVENTOR(S):

Sumitomo Chemical Company, Limited, Japan PATENT ASSIGNEE(S):

SOURCE: PCT Int. Appl., 47 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	·DATE
WO 2004046125	A1	20040603	WO 2003-JP13750	20031028
W: AE, AG,	AL, AM, AT	, AU, AZ,	BA, BB, BG, BR, BY, BZ,	CA, CH, CN,
CO. CR.	CU. CZ. DE	. DK. DM.	DZ. EC. EE. EG. ES. FI.	GB, GD, GE,

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GH, GM, HR, HU, ID, IL, IN, IS, KE, KG, KR, KZ, LC, LK, LR, LS,
             LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG,
             PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR,
             TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
             FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,
             BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
    AU 2003274762
                                            AU 2003-274762
                          A1
                                20040615
                                                                    20031028
                                            EP 2003-758967
    EP 1569922
                                20050907
                          Α1
                                                                    20031028
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
    BR 2003016439
                                            BR 2003-16439
                          Α
                                20051011
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    CN 1714086
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                                20051228
                                            CN 2003-80103741
                                                                    20031028
    JP 2004182722
                          Α
                                20040702
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                                20060309
                                            US 2005-532478
                                                                    20050422
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                                20071127
    ZA 2005003460
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                                20060830
                                            ZA 2005-3460
                                                                    20050429
    MX 2005PA05336
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                                20050725
                                            MX 2005-PA5336
                                                                    20050518
    IN 2005CN00988
                                20070622
                                            IN 2005-CN988
                                                                    20050520
PRIORITY APPLN. INFO.:
                                            JP 2002-337884
                                                                    20021121
                                                                 Α
                                            WO 2003-JP13750
                                                                 W
                                                                    20031028
```

OTHER SOURCE(S):

MARPAT 141:23538

GI

$$N-S$$
 $R^{1}O$
 N
 $A^{1}R^{2}$
 I
 $H_{3}C$
 I

AB The invention relates to novel 1,2,4-thiadiazole compds. I [wherein: R1 = C3-7 alkynyl that may be substituted with halo; R2 = C3-8 cycloalkyl which may be substituted with C1-4 alkyl, halo, CF3, or the like; A1 = bond, C1-2 alkylene, or C2-3 alkylidene]. I have excellent arthropod-controlling activity, and can effectively control arthropod pests such as insects, acarids, and the like. Examples include 15 product syntheses, 8 precursor prepns., 8 formulations, and 1 bioassay. For instance, 5-chloro-3-(methylthio)-1,2,4-thiadiazole was alkylated with cyclohexylzinc bromide using a Pd complex catalyst, and Soxidized using m-CPBA, to give 3-(methylsulfonyl)-5-cyclohexyl-1,2,4thiadiazole. This sulfone was coupled with 2-butyn-1-ol using NaH in DMF, to give invention compound 5-cyclohexyl-3-(2-butynyloxy)-1,2,4- thiadiazole (II). At 500 ppm (spray) against Aphis gossypii on cucumber seedlings, each exemplified compound I reduced the number of plant parasites from 20 individuals to not greater than 3.

L13 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2004:333715 CAPLUS Full-text

DOCUMENT NUMBER: 140:339331

TITLE: Preparation of thiadiazole compounds as

arthropodicides

INVENTOR(S): Ihara, Hideki

Sumitomo Chemical Company, Limited, Japan PATENT ASSIGNEE(S):

SOURCE: PCT Int. Appl., 70 pp.

CODEN: PIXXD2

DOCUMENT TYPE: LANGUAGE:

Patent Japanese

FAMILY ACC. NUM. COUNT:

1 1

PATENT INFORMATION:

PA	TENT :	NO.			KIN	D	DATE			APPI	LICAT	ION :	NO.		D	ATE	
WO	2004	0334	52		A1	-	2004	0422		WO 2	2003-	JP12	831		2	0031	007
	W:	ΑE,	AG,	AL,	AM,	AT,	ΑU,	ΑZ,	BA,	BB,	, BG,	BR,	BY,	ΒZ,	CA,	CH,	CN,
		CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	, EE,	EG,	ES,	FI,	GB,	GD,	GE,
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		PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	, SK,	SL,	SY,	ТJ,	TM,	TN,	TR,
		TT,	TZ,	UA,	ŪG,	US,	UZ,	VC,	VN,	YU,	, ZA,	ZM,	zw				
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		KG,	ΚZ,	MD,	RU,	ТJ,	TM,	ΑT,	BE,	BG,	, CH,	CY,	CZ,	DE,	DK,	EE,	ES,
		FI,	FR,	GB,	GR,	HU,	IE,	IT,	LU,	MC,	, NL,	PT,	RO,	SE,	SI,	SK,	TR,
		BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	, GW,	ML,	MR,	NE,	SN,	TD,	TG
JP	2004	1314	38		Α		2004	0430		JP 2	2002-	2984	89		2	0021	011
AU	2003	2711	12		A1		2004	0504		AU 2	2003-	2711	12		2	0031	007
EP	1550	661			A1		2005	0706		EP 2	2003-	7513	59		2	0031	007
	R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	, IT,	LI,	LU,	NL,	SE,	MC,	PT,
		IE,	SI,	LT,	LV,	FI,	RO,	MK,	CY,	AL,	TR,	BG,	CZ,	EE,	HU,	SK	
BR	2003	0146	16		Α		2005	0726		BR 2	2003-	1461	6		2	0031	007
	1703				Α		2005	1130		CN 2	2003-	8010					
ZA	2005	0024	99		Α		2006	0628		ZA 2	2005-	2499			2	0031	007
US	2006	0149	62		A1		2006	0119		US 2	2005-	5283	98		2	0050	317
MX	2005	PA03	666		Α		2005	0608		MX 2	2005-	PA36	66		2	0050	406
IN	2005	CN00	575		Α		2007	0622		IN 2	2005-	CN57	5		2	0050	407
RIORIT	Y APP	LN.	INFO	.:						JP 2	2002-	2984	89	7	A 2	0021	011
										WO 2	2003-	JP12	831	1	₩ 2	0031	007
THER SO	OURCE	(S):			MAR	PAT	140:	3393	31								

AB Title compds. I (R1 = alkyl, alkenyl, alkynyl, etc.; R2 = heterocyclylalkyl) are prepared Thus, reaction of 5-chloro-3-(4-methylbenzyl)thio-1,2,4-thiadiazole with 2,2-dimethyl-1,3-dioxolane-4-methanol in DMF in the presence of NaH at room temperature for 4 h gave 5-(2,2-dimethyl-1,3-dioxolan-4-yl)methoxy-3-(4-methylbenzyl)thio-1,2,4-thiadiazole (II). II showed arthropodicidal activity against Aphis gossypii at 500 ppm.

REFERENCE COUNT:

THERE ARE 14 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2007 ACS on STN ACCESSION NUMBER: 2003:570972 CAPLUS Full-text DOCUMENT NUMBER: 139:133570

14

TITLE:

Preparation of 5-alkyloxy-3-alkylthio-1,2,4-

thiadiazole derivatives having control activities against injurious arthropods

INVENTOR(S):

Ihara, Hideki; Sakamoto, Noriyasu; Tomioka,

Hiroki

PATENT ASSIGNEE(S):

Sumitomo Chemical Company, Limited, Japan

SOURCE:

LANGUAGE:

PCT Int. Appl., 65 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PA'	rent :				KIN								NO.			DATE	
WO	2003												- 37			20030	115
	W:	ΑE,	AG,	AL,	AM,	ΑT,	ΑU,	ΑZ,	BA,	BE	3, BO	, BR	, BY,	ΒZ,	CA	, сн,	CN,
													, FI,				
													, LC,				
		LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	ΜX	, M2	, NO	, NZ,	OM,	PH	, PL,	PT,
		RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	ТJ	, TN	I, TN	, TR,	TT,	TZ	, UA,	UG,
		US,	UZ,	VC,	VN,	ΥU,	ZA,	ZM,	zw								
	RW:	GH,	GM,	ΚE,	LS,	MW,	MZ,	SD,	SL,	SZ	, T2	, UG	, ZM,	ZW,	AM	, AZ,	BY,
													, CZ,				
		FI,	FR,	GB,	GR,	HU,	ΙE,	IT,	LU,	MC	, NI	, PT	, SE,	SI,	SK	, TR,	BF,
		ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW	, MI	, MR	, NE,	SN,	TD	, TG	
AU	2003	2032	24		A1		2003	0730		AU	2003	-203	224			20030	115
JP	2003	2773	72		Α		2003	1002		JΡ	2003	-674	6			20030	115
EP	1475	374			A1		2004	1110		ΕP	2003	-701	718			20030	115
	R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GF	R, IT	, LI	, LU,	NL,	SE	, MC,	PT,
		ΙE,	SI,	LT,	LV,	FI,	RO,	MK,	CY,	ΆI	, TF	, BG	, CZ,	EE,	HU	, SK	
BR	2003	0068	23		Α		2004	1221		BR	2003	-682	3			20030	115
CN	1617	863			Α		2005	0518		CN	2003	-802	385			20030	115
US	2005	2155	78		A1		2005	0929		US	2004	-498	651			20040	610
	7273						2007										
IN	2004	CN01	534		Α		2006	0210		IN	2004	-CN1	534			20040	709
US	2007	2935	10		A1		2007	1220		US	2007	-880	846			20070	724
PRIORITY	Y APP	LN.	INFO	.:						JP	2002	-835	6		A	20020	117
										WO	2003	-JP2	37	,	W	20030	115
										US	2004	-498	651		A 3	20040	610
OTHER SO	OURCE	(S):			MAR	PAT	139:	13357	70								

$$\begin{array}{c} \text{MeS} \\ \text{N} \\ \text{N} \\ \text{S} \\ \text{OR2} \quad \text{I} \end{array}$$

16

AB The title thiadiazole compds. with general formula of I, which have excellent control activities against injurious arthropods, [wherein R1 = Me, alkenyl, alkoxyalkyl, alkylthioalkyl, alkoxyalkoxyalkyl, alkylthioalkoxyalkyl, (un) substituted PhO-alkyl, Ph-alkoxyalkyl, or acyloxyalkyl; R2 = (un) substituted Ph-alkyl, Py-alkyl, or pyrimidylalkyl] are prepared For example, 5-chloro-3-methylthio-1,2,4-thiadiazole was reacted with PhCH2OH in DMF in the presence of NaH to give 5-benzyloxy-3-methylthio-1,2,4-thiadiazole (II). I at concentration of 500 ppm killed more than 85% of aphis gossypii on cucumber seedlings in 6 days.

REFERENCE COUNT:

THERE ARE 16 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L13 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2002:900807 CAPLUS Full-text

DOCUMENT NUMBER:

137:381259

TITLE:

Preparation of 1,2,4-thiadiazole compounds

and arthropodicides containing them Ihara, Hideki; Sakamoto, Noriyasu

PATENT ASSIGNEE(S):

Sumitomo Chemical Co., Ltd., Japan

SOURCE:

Jpn. Kokai Tokkyo Koho, 13 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

INVENTOR(S):

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PA'	TENT 1	NO.			KIN	D	DATE				ICAT				Γ	ATE	
JP	2002	3385	57		A	_	2002	1127							2	0010	- 522
WO	2004	0417	98		A1		2004	0521		WO 2	002-	JP11	644		2	0021	108
	W:	ΑE,	AG,	AL,	AM,	AT,	ΑU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	BZ,	CA,	CH,	CN,
		CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GE,	GH,
							IN,										
		LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NO,	NZ,	OM,	PH,	PL,	PT,
		RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	ТJ,	TM,	TN,	TR,	TT,	TZ,	UA,	UG,
		US,	UZ,	VN,	YU,	ZA,	ZM,	ZW`									
	RW:	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	ŪG,	ZM,	ZW,	AM,	ΑZ,	BY,
		KG,	ΚZ,	MD,	RU,	ТJ,	TM,	ΑT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,
		FI,	FR,	GB,	GR,	ΙE,	IT,	LU,	MC,	NL,	PT,	SE,	SK,	TR,	BF,	ВJ,	CF,
		CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,	NE,	SN∙,	TD,	TG			
AU	2002	3683	30		A1		2004	0607		AU 2	002-	3683	30		2	0021	108
BR	2002	0159	11		Α		2005	0726		BR 2	002-	1591	1		2	0021	108
EP	1574	505			A1		2005	0914		EP 2	002-	8081	00		2	0021	108
	R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	PT,
		IE,	SI,	LT,	LV,	FI,	RO,	MK,	CY,	AL,	TR,	BG,	CZ,	EE,	SK		
	1688				À		2005										
US	2006	1672	66		A1		2006	0727		US 2	005-	5301	36		2	0050	404
ZA	2005	0029	10		Α		2006	0628		ZA 2	005-	2910			2	0050	411
MX	2005	PA04	824		Α		2005	0722		MX 2	005-3	PA48	24		2	0050	504
IN	2005	CN00	857		Α		2007	0810		IN 2	005-0	CN85	7	•	2	0050	506
PRIORIT	Y APP	LN.	INFO	.:						JP 2	001-	1522	69		A 2	0010	522
										WO 2	002-	JP11	644		A 2	0021	108
OTHER SO	OURCE	(S):			MAR	PAT	137:	3812	59								

$$R^{10}$$
 R^{2} R^{2}

AΒ The compds. I [R1 = C3-7 (halo)alkenyl; R2 = halo, C1-4 alkyl, C1-3 haloalkyl, C1-4 haloalkoxy, cyano, NO2; n = 0-5; A = 0, S, direct bond, CR3R4, NR5; R3, R4 = H, C1-4 alkyl; R5 = H, C1-7 alkyl, C1-3 haloalkyl, C2-4 (halo)alkoxyalkyl, C3-6 (halo)alkenyl, C3-7 (halo)alkynyl, CH2CN] and arthropod control agents containing I are claimed. A composition containing 5-phenyl-3-propargyloxy1,2,4-thiadiazole (preparation given), showed ≥90% control against Aphis gossypii parasitic on cucumber seedlings.

L13 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

1997:481831 CAPLUS Full-text

DOCUMENT NUMBER:

127:205546

TITLE:

Studies of pyrazines. Part 33. Synthesis of 2,3-diaminopyrazines via [1,2,5]thiadiazolo

[3,4-b]pyrazines

AUTHOR(S):

Sato, Nobuhiro; Mizuno, Hajime

CORPORATE SOURCE:

Department Chemistry, Yokohama City University,

Yokohama, 236, Japan

SOURCE:

Journal of Chemical Research, Synopses (1997), (7),

250-251

CODEN: JRPSDC; ISSN: 0308-2342

PUBLISHER:

Royal Society of Chemistry

DOCUMENT TYPE:

Journal

LANGUAGE:

English

OTHER SOURCE(S):

CASREACT 127:205546

GI

$$R^1$$
 N N S R^2 N N N N

AB The syntheses of [1,2,5]thiadiazolo[3,4-b]pyrazines I (R1 = H, Me, Ph, R2 = H; R1 = Me, R2 = H, Me; R1 = Ph, R2 = Me; R1 = R2 = Ph) as well as their reduction to 2,3-diaminopyrazines II are described.

REFERENCE COUNT:

THERE ARE 15 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d his nofil

(FILE 'HOME' ENTERED AT 15:07:22 ON 28 DEC 2007)

FILE 'REGISTRY' ENTERED AT 15:07:56 ON 28 DEC 2007

L1 STR

L2 3 SEA SSS SAM L1

L3 · 98 SEA SSS FUL L1

FILE 'CAPLUS' ENTERED AT 15:09:48 ON 28 DEC 2007

L4 21 SEA ABB=ON PLU=ON L3

E IHARA H/AU

L5 242 SEA ABB=ON PLU=ON ("IHARA H"/AU OR "IHARA HIDEAKI"/AU OR

"IHARA HIDEKI"/AU)

E TAKAOKA D/AU

L6 51 SEA ABB=ON PLU=ON ("TAKAOKA D"/AU OR "TAKAOKA DAISUKE"/AU)

E MIZUNO H/AU

L7 186 SEA ABB=ON PLU=ON ("MIZUNO H"/AU OR "MIZUNO HAJIME"/AU)

L8 474 SEA ABB=ON PLU=ON (L5 OR L6 OR L7)

L9 6 SEA ABB=ON PLU=ON L8 AND THIA? AND ?AZOL?

L10 1 SEA ABB=ON PLU=ON L9 AND L4

L11	1	SEA ABB=ON	PLU=ON	L4 AND L8
L12	21	SEA ABB=ON	PLU=ON	L4 OR L10
L13	5	SEA ABB=ON	PLU=ON	L9 NOT L10

FILE 'CAPLUS' ENTERED AT 15:12:41 ON 28 DEC 2007

D QUE L12

D L12 IBIB ABS HITSTR TOT

D QUE L13

D L13 IBIB ABS TOT